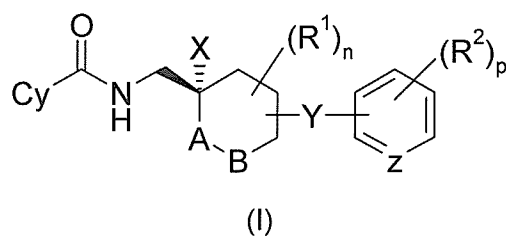


Amendments to the Abstract:

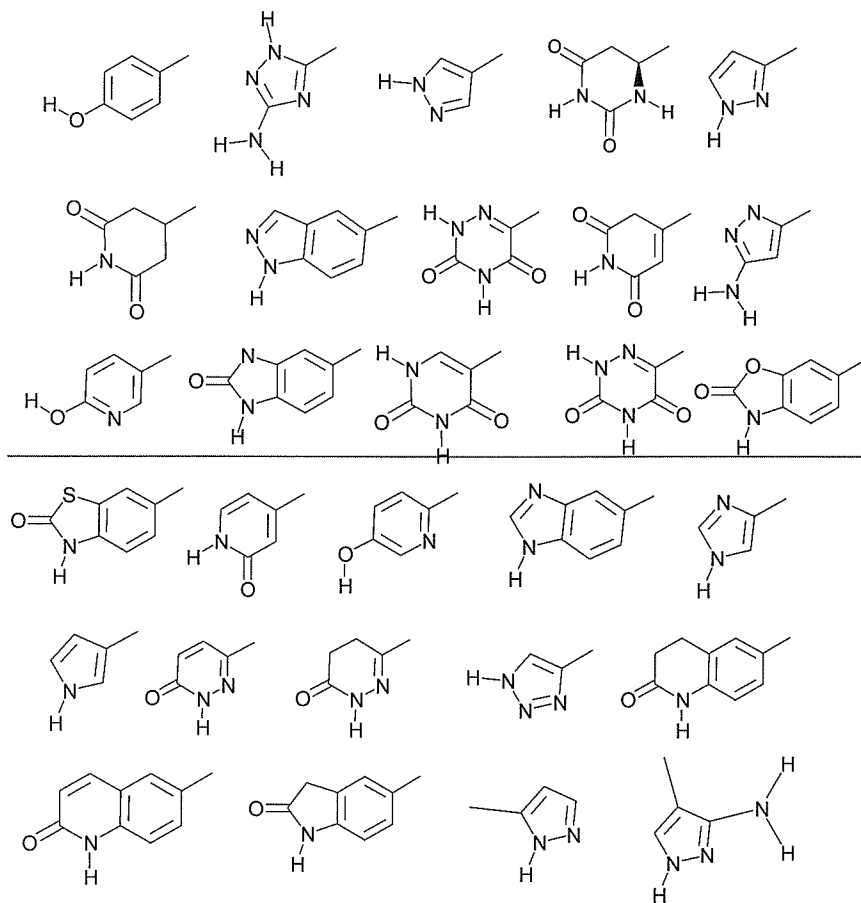
The present invention relates to compounds of the formula (I):



or a pharmaceutically acceptable salt or solvate thereof, wherein:

A and B independently represent CH_2 or O, with the proviso that A and B are not simultaneously O;

Cy represents one of the following



~~optionally substituted by one to three groups selected from hydroxy, halogen, C₁₋₆alkyl, C₁₋₆alkoxy, C₁₋₆haloalkyl, C₁₋₆alkylamino and amino;~~

~~R¹ and R² are independently selected from hydroxy, halogen, C₁₋₆alkyl, C₁₋₆alkoxy, C₁₋₆haloalkyl and C₃₋₈cycloalkyl;~~

~~n represents an integer from 0-4;~~

~~X is hydrogen, hydroxy, halogen or C₁₋₆alkoxy;~~

~~Y is oxy, thio, a 1-4 membered alkylene, a 2-4 membered alkylene ether, 2-4 membered alkylene thioether or an oxyethyleneoxy group, optionally substituted by 1 to 4 groups independently selected from hydroxy, halogen, C₁₋₆alkyl, C₁₋₆alkoxy and C₁₋₆haloalkyl;~~

~~Z is CH or N; and~~

~~p represents an integer from 0-5 when Z is CH or 0-4 when Z is N; when p represents 2 or more, two of R²s may be taken together with the carbon atoms to which they are attached to form a 5-8 membered cycloalkyl ring~~

~~to processes for the preparation of, intermediates used in the preparation of, compositions containing such compounds and the uses of such compounds as antagonists of the NMDA NR2B receptor.~~